



MEDI-CAL UPDATE

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www.medi-cal.ca.gov

Billing and Policy Pharmacy Bulletin 567

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The energy challenge facing California is real. The Department of Health Services encourages practical and feasible energy saving measures while considering the health and safety of clients, workers and family members.

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Medi-Cal List of Contract Drugs: Update

The following provider manual section has been updated: *Drugs: Contract Drugs List Part 1 – Prescription Drugs.*

Changes, effective October 1, 2003

<u>Drug</u>	<u>Size and/or Strength</u>	<u>Billing Unit</u>
‡ * ABACAVIR SULFATE		
Tablets	300 mg	ea
Liquid	20 mg/cc	cc
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		
‡ * ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE		
Tablets	300 mg/150 mg/300 mg	ea
* Restricted to use alone or as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		
‡ * AMPRENAVIR		
Capsules	50 mg	ea
	150 mg	ea
Oral solution	15 mg/cc	cc
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		

Please see **Contract Drugs**, page 3

EDS/MEDI-CAL HOTLINES

Border Providers	(916) 636-1000, ext. 2100
Computer Media Claims (CMC).....	(916) 636-1100
DHS Medi-Cal Fraud Hotline.....	1-800-822-6222
Health Access Programs (HAP) – OB, CPSP, Family PACT, BCEDP Providers.....	1-800-257-6900
POS/Internet Help Desk	1-800-427-1295
Provider Support Center (PSC).....	1-800-541-5555
Provider Telecommunications Network (PTN).....	1-800-786-4346
Specialty Programs	1-800-541-7747

For a complete listing of specialty programs and hours of operation, please refer to the Medi-Cal Directory in the provider manual.

**MEDI-CAL FRAUD
IS AGAINST THE
LAW**

**MEDI-CAL FRAUD COSTS TAXPAYERS MILLIONS
EACH YEAR AND CAN ENDANGER
THE HEALTH OF CALIFORNIANS.**

**HELP PROTECT MEDI-CAL AND YOURSELF
BY REPORTING YOUR OBSERVATIONS TODAY.**

**DHS MEDI-CAL FRAUD HOTLINE
1-800-822-6222**

THE CALL IS FREE AND YOU CAN REMAIN ANONYMOUS.

Knowingly participating in fraudulent activities can result in prosecution and jail time. Help prevent Medi-Cal fraud.

Contract Drugs (continued)

Changes, effective October 1, 2003

<u>Drug</u>	<u>Size and/or Strength</u>	<u>Billing Unit</u>
‡ * DELAVIRIDINE MESYLATE		
Tablets	100 mg	ea
	200 mg	ea
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection.</u>		
‡ * DIDANOSINE		
Capsules, delayed release, E.C.	125 mg	ea
	200 mg	ea
	250 mg	ea
	400 mg	ea
Tablets	25 mg	ea
	50 mg	ea
	100 mg	ea
	150 mg	ea
	200 mg	ea
Powder for oral solution	100 mg/packet	ea
	167 mg/packet	ea
	250 mg/packet	ea
	375 mg/packet	ea
Pediatric powder for oral solution	20 mg/cc	cc
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection.</u>		
‡ * EFAVIRENZ		
Capsules	50 mg	ea
	100 mg	ea
	200 mg	ea
Tablets	600 mg	ea
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection.</u>		
FENOFIBRATE		
Tablets	54 mg	ea
	160 mg	ea
<u>(NDC labeler code 00074 [Abbott Laboratories] only.)</u>		
FENOFIBRATE, MICRONIZED		
Capsules	67 mg	ea
	134 mg	ea
	200 mg	ea
<u>(NDC labeler code 00074 [Abbott Laboratories] only.)</u>		

Please see Contract Drugs, page 4

Contract Drugs (continued)

Changes, effective October 1, 2003

<u>Drug</u>	<u>Size and/or Strength</u>	<u>Billing Unit</u>
‡ * FLUCONAZOLE		
Injection	2 mg/cc	100 cc (saline)
		200 cc (saline)
		100 cc (dextrose)
		200 cc (dextrose)
Tablets	50 mg	ea
	100 mg	ea
	150 mg	ea
	200 mg	ea
* Restricted to use in cancer patients, in patients with AIDS or an AIDS-related condition, in the treatment of vaginal candidiasis, and in the treatment of infections caused by <i>Coccidioides immitis</i> Human Immunodeficiency Virus (HIV) infection.		
‡ * INDINAVIR SULFATE		
Capsules	100 mg	ea
	200 mg	ea
	333 mg	ea
	400 mg	ea
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		
‡ * LAMIVUDINE		
Tablets	150 mg	ea
	300 mg	ea
Liquid	10 mg/cc	cc
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		
‡ * LAMIVUDINE AND ZIDOVUDINE		
Tablets	150 mg/300 mg	ea
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		
‡ * LOPINAVIR AND RITONAVIR		
Capsules	133.3 mg – 33.3 mg	ea
Oral solution	400 mg – 100 mg/5 cc	cc
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection.		
LOXAPINE HCL		
Solution	25 mg/cc	cc
Injection	50 mg/cc	cc
(Loxitane by Watson Laboratories [NDC labeler code 52544] only.)		
LOXAPINE SUCCINATE		
Capsules	5 mg	ea
	10 mg	ea
	25 mg	ea
	50 mg	ea
(Loxitane by Watson Laboratories [NDC labeler code 52544] only.)		

Please see Contract Drugs, page 5

Contract Drugs (continued)

Changes, effective October 1, 2003

<u>Drug</u>	<u>Size and/or Strength</u>	<u>Billing Unit</u>
‡ * NELFINAVIR MESYLATE		
Tablets	250 mg	ea
Oral powder	50 mg/Gm	Gm
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * NEVIRAPINE		
Tablets	200 mg	ea
Liquid	50 mg/5cc	cc
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * RITONAVIR		
Capsules	100 mg	ea
Solution	80 mg/cc	cc
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * SAQUINAVIR		
Capsules	200 mg	ea
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * SAQUINAVIR MESYLATE		
Capsules	200 mg	ea
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * STAVUDINE		
Capsules	15 mg	ea
	20 mg	ea
	30 mg	ea
	40 mg	ea
Powder for oral solution	1 mg/cc	cc
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * TENOFOVIR DISOPROXIL FUMARATE		
Tablets	300 mg	ea
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		
‡ * ZALCITABINE		
Tablets	0.375 mg	ea
	0.750 mg	ea
* Restricted to use <u>as combination therapy</u> in the treatment of AIDS or an AIDS-related condition <u>Human Immunodeficiency Virus (HIV) infection</u> .		

Please see Contract Drugs, page 6

Contract Drugs (*continued*)**Changes, effective October 1, 2003**

<u>Drug</u>	<u>Size and/or Strength</u>	<u>Billing Unit</u>
‡ * ZIDOVUDINE		
Tablets	300 mg	ea
Capsules	100 mg	ea
Liquid	50 mg/5cc	cc
Injection	10 mg/cc	cc
* Restricted to use as combination therapy in the treatment of AIDS or an AIDS-related condition Human Immunodeficiency Virus (HIV) infection .		

Changes, effective November 1, 2003

MOEXIPRIL HCL		
+ Tablets	7.5 mg	ea
	15 mg	ea
<u>(NDC labeler code 00091 [Schwarz Pharma, Inc.] only.)</u>		
NIACIN		
Tablets, extended release	500 mg	ea
	750 mg	ea
	1000 mg	ea
<u>(NDC labeler code 60598 [KOS Pharmaceuticals, Inc.] only.)</u>		
PAROXETINE HCL		
Suspension, oral	10 mg/5cc	cc
Tablets	10 mg	ea
	20 mg	ea
	30 mg	ea
	40 mg	ea
Tablets, controlled release	12.5 mg	ea
	25 mg	ea
	37.5 mg	ea
<u>(NDC labeler code 00029 [SmithKline Beecham] only.)</u>		

‡ Drug is exempt from the monthly drug claim limit line.

+ Frequency of billing requirement.

Refer to manual replacement pages drugs cdl p1a 1, 10, 37 and 41 (Part 2), drugs cdl p1b 1, 10, 12, 28, 35, 42, 43 and 53 (Part 2), drugs cdl p1c 2, 3, 15 and 35 (Part 2) drugs cdl p1d 1, 4, 8, 22 and 23 (Part 2).

Authorized Drug Manufacturer Labeler Codes: Update

The *Drugs: Contract Drugs List Part 5 – Authorized Drug Manufacturer Labeler Codes* section has been updated as follows.

Additions, effective October 1, 2003

NDC Labeler Code	Contracting Company's Name
67871	QOL MEDICAL
68013	VISION PHARMA, LLC

Additions, effective January 1, 2004

NDC Labeler Code	Contracting Company's Name
68032	RIVER'S EDGE PHARMACEUTICALS
67386	OVATION PHARMACEUTICALS, INC.
68134	PALMETTO PHARMACEUTICALS, INC.

Reinstatements, effective October 1, 2003

NDC Labeler Code	Contracting Company's Name
00463	C.O. TRUXTON, INC.

Terminations, effective January 1, 2004

NDC Labeler Code	Contracting Company's Name
59229	HORUS THERAPEUTICS

These updates are reflected on manual replacement pages drugs cdl p5 4, 10, 14 and 15 (Part 2).

Compound Drug Policies: Update

Effective for dates of service on or after September 22, 2003, the following policy changes are enacted:

Code I Changed for 10-day After Discharge Coverage of Unlisted Intravenous or Intra-arterial Drugs

The Code I restriction on Parenteral Nutrition Solutions (TPN or Hyperalimentation) and Separately Administered Intravenous Lipids is being standardized to match Code I restrictions for Intravenous Solutions of Unlisted Antibiotics and Intravenous Solutions of Other Unlisted Drugs. The new Code I restriction reads:

“Restricted to dispensing within 10 days following inpatient discharge from an acute care hospital, when I.V. therapy with the same product was started before discharge. There is a maximum of 10 days supply per dispensing within the 10-day period.”

Creating a standardized 10-day period is expected to improve the consistency of claim payments.

Note: Non-compounded products must be billed using the product's NDC number and must be billed using the *Pharmacy Claim Form* (30-1). Non-compounded intravenous products not listed on the *Medi-Cal List of Contract Drugs* continue to require a *Treatment Authorization Request* (TAR) even if dispensed during the 10-day post-discharge window.

Please see Compound, page 8

Compound (*continued*)**Container Count Restriction, Single-Ingredient Injections**

Single-ingredient injections (sterile transfers) for more than seven containers require a TAR, regardless of the 10-day post-discharge window or *Medi-Cal List of Contract Drugs* status.

Container Count Restrictions, Multiple-Ingredient Injections (more than 20 containers)

Multiple-ingredient injections for more than 20 containers require a TAR, regardless of the 10-day post-discharge window or *Medi-Cal List of Contract Drugs* status.

Enforcement of List of Contract Drugs for Ingredients in Compounded Drugs

All ingredients contained in a compounded product must be listed on the *Medi-Cal List of Contract Drugs*. If one or more ingredients is not on the list, the product requires a TAR.

Inactive Ingredients in Compound Drugs

The Department of Health Services (DHS) recognizes that certain “pharmaceutic aids” not on the *Medi-Cal List of Contract Drugs* are required to manufacture compound drugs. These might include vehicles, adjuvants, suspending, flavoring and coloring agents, etc. A TAR generally is not required for these types of ingredients when included in a compound drug. To determine if an inactive ingredient requires a TAR, submit the claim for the compound online using Real-Time Internet Pharmacy (RTIP) claim submission or the National Council for Prescription Drug Programs (NCPDP)-compliant software. If the claim denies, the online denial message will note which ingredients caused the denial.

Non-Compound Products Billing

Non-compounded products must be billed using the product’s NDC number and must be billed using the *Pharmacy Claim Form* (30-1).

Effective Date for Claims Filed on the Pharmacy Claim Form (30-1)

The above changes are not effective until November 1, 2003 for claims submitted using the old compound drug format (for example, submitted on the *Pharmacy Claim Form* [30-1] with attached compounding sheet). The delay allows pharmacies a window in which they can accommodate recipients who began treatment before notification of the changes.

The updated information is reflected on manual replacement pages drugs cdl p1b 30 and 31 (Part 2) and iv sol spec 2 thru 4 (Part 2).

Billing Code 9999A: Change in Billing Instructions

Effective November 1, 2003, providers billing the 9999A code for Unlisted and Miscellaneous Medical Services must submit a copy of the original Treatment Authorization Request (TAR) along with appropriate pricing documentation (for example, invoice or manufacturer catalog page) with the claim. *The updated information is reflected on manual replacement pages mc sup intro 3 (Part 2), mc sup lst4 11 (Part 2), pcf30-1 comp 11 (Part 2) and tar comp 10 (Part 2).*

Instructions for Manual Replacement Pages

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Part 2

Remove and replace:

drugs cdl p1a 1/2, 9/10, 37/38 and 41/42
drugs cdl p1b 1/2, 9 thru 12, 27 thru 32, 35/36, 41 thru 44 and 53/54
drugs cdl p1c 1 thru 4, 15/16 and 35/36
drugs cdl p1d 1 thru 4, 7/8 and 21 thru 23
drugs cdl p5 3/4, 9/10 and 13 thru 15
iv sol spec 1 thru 4
mc sup intro 3
mc sup lst4 11
pcf30-1 comp 11/12
tar comp 9/10